AMENDMENTS TO THE SPECIFICATION

Please amend the paragraph beginning on line 10, page 68 as follows:

Cytokine specific primer pairs for the inducible MxA mRNA and the constitutively expressed G3PHD mRNA were designed by JSW and otained from GynSys, the Woodlands, TX. The primer pairs are complimentary complementary to exons separated by at least one intron to avoid unrecognized amplification of cellular DNA and are designated by the relative position in the human mRNA {Mx: Mx: Plus strand primer GTGGAGCAGGACCTGGCCCTG (400-420) (SEO_ID_NO: 1); minus strand primer GAGCCTCTGTGGTGGCAATG (895-876) (SEQ_ID_NO: 2); G3PDH: G3PDH: plus strand primer CAACGGATTTGGTCGTATTGGGCGC (84-108) (SEO ID NO: 3); minus strand primer TTACTCCTTGGAGGCCATGTGGGCC (1068-1094) (SEQ ID NO: 4)}. 5 µl of cDNA (representing DNA derived from 3-5 x 10⁶ cells) was amplified in 0.5 ml Gene-amp reaction tubes (Cetus Corp) in 200 µM final concentration of the four primers, 200 mM dNTPs, 0.5 U of Taq polymerase, and PCR buffer containing 2.5 mM MgCl₂, 50 mM KCL KCl, 10 Mm mM Tris-Cl pH 8.3, and 0.001% gelatin in a final volume of 25 ml. The reaction mixture was overlaid with a drop oflight mineral oil, and PCR performed in a DNA thermal cycler (M.J. Research) for 30 cycles: 60 seconds denaturation at 94°C, 120 seconds annealing at

60°C, and 3 minutes extension at 72°C. The reaction product was visualized by electrophoresis of 20 µl of the reaction mixture at 80 V for 70 minutes in 2% agarose in 0.5x TBE buffer containing 0.5 mg/ml ethidium bromide. Specificity of the amplified product was validated by the predicted size following transfer to nitrocellulose, hybridization with 3'-digoxigenin-labeled (Genius(5, Boeringer Mannheim, Indianapolis, IN) internal oligonucleotide probes (MxA: ACCAGATGCCCTCTGGTGCTG, 405-425 (SEQ_ID_NO: 5); and GAPDH: CCGTCTCCAGAACATCATCCCTGCC, 687-663 (SEO ID NO: 6), and chemiluminescent detection with an alkaline phosphatase-conjugated anti-digoxigenin antibody and [4-methoxy-4-(3phosphatephenyl)spiro(1,2-dioxetane(3,2'-adamantane)] substrate (Genius (7; Boeringer Mannheim) and exposure for 15 and 30 minutes to X-ray film. The relative amount of MxA and GAPDH message amplified was determined by densitometric analysis of the bands on exposed film.